

- Ant A3* 31. 42. The method according to claim 39, wherein said modulator is an antibody.
- 32. 43.* The method according to claim 41, wherein said polypeptide is provided as part of a membrane fraction.
- 33. 44.* The method according to claim 39, wherein said HIV virus is HIV-1 or HIV-2
- 34. 45.* The method according to claim 39, wherein said HIV virus is a macrophage-trophic (M trophic or R5) strain virus.
- 35. 46.* The method according to claim 39, wherein said decreasing of infectivity is monitored by measuring a modification of the signaling activity of said CCR5 chemokine receptor.
- 36. 47.* The method according to claim 46, wherein said measuring of signaling activity comprises measuring of one or more of: changes in levels of cellular acidification, intracellular calcium, IP<sub>3</sub>, and stimulation of an intracellular cascade.
- 37. 48.* The method according to claim 39, wherein said decreasing of infectivity is monitored by measuring production of an HIV polypeptide.
- 38. 49.* The method according to claim 48, wherein said HIV polypeptide is p24.
- 39. 50.* A CCR5 chemokine receptor modulator which decreases the infectivity of a cell expressing said CCR5 chemokine receptor by at least two-fold when delivered to said cell.
- 40. 51.* The CCR5 chemokine receptor modulator according to claim 50, wherein said modulator is an antibody.

#### REMARKS

Upon entry of this amendment, claims 39 to 51 are pending. No new matter is introduced